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NEWS 4 AUG 28 ADISCTI Reloaded and Enhanced
NEWS 5 AUG 30 CA(SM)/CAplus(SM) Austrian patent law changes
NEWS 6 SEP 21 CA/CAplus fields enhanced with simultaneous left and right truncation
NEWS 7 SEP 25 CA(SM)/CAplus(SM) display of CA Lexicon enhanced
NEWS 8 SEP 25 CAS REGISTRY(SM) no longer includes Concord 3D coordinates
NEWS 9 SEP 25 CAS REGISTRY(SM) updated with amino acid codes for pyrrolysine
NEWS 10 SEP 28 CEABA-VTB classification code fields reloaded with new classification scheme
NEWS 11 OCT 19 LOGOFF HOLD duration extended to 120 minutes
NEWS 12 OCT 19 E-mail format enhanced
NEWS 13 OCT 23 Option to turn off MARPAT highlighting enhancements available
NEWS 14 OCT 23 CAS Registry Number crossover limit increased to 300,000 in multiple databases
NEWS 15 OCT 23 The Derwent World Patents Index suite of databases on STN has been enhanced and reloaded
NEWS 16 OCT 30 CHEMLIST enhanced with new search and display field
NEWS 17 NOV 03 JAPIO enhanced with IPC 8 features and functionality
NEWS 18 NOV 10 CA/CAplus F-Term thesaurus enhanced
NEWS 19 NOV 10 STN Express with Discover! free maintenance release Version 8.01c now available
NEWS 20 NOV 20 CAS Registry Number crossover limit increased to 300,000 in additional databases
NEWS 21 NOV 20 CA/CAplus to MARPAT accession number crossover limit increased to 50,000
NEWS 22 DEC 01 CAS REGISTRY updated with new ambiguity codes
NEWS 23 DEC 11 CAS REGISTRY chemical nomenclature enhanced
NEWS 24 DEC 14 WPIDS/WPINDEX/WPIX manual codes updated
NEWS 25 DEC 14 GBFULL and FRFULL enhanced with IPC 8 features and functionality
NEWS 26 DEC 18 CA/CAplus pre-1967 chemical substance index entries enhanced with preparation role
NEWS 27 DEC 18 CA/CAplus patent kind codes updated
NEWS 28 DEC 18 MARPAT to CA/CAplus accession number crossover limit increased to 50,000
NEWS 29 DEC 18 MEDLINE updated in preparation for 2007 reload
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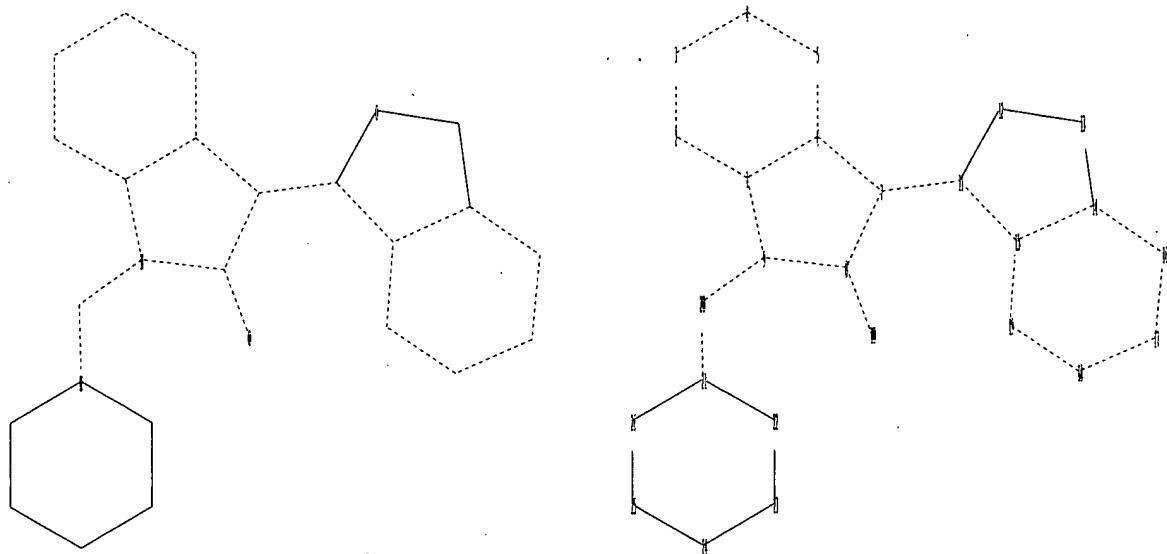
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<http://www.cas.org/ONLINE/UG/reqprops.html>

=>
Uploading C:\Program Files\Stnexp\Queries\10829139.str

10829139



chain nodes :

10 20

ring nodes :

1 2 3 4 5 6 7 8 9 11 12 13 14 15 16 17 18 19 21 22 23 24 25
26

chain bonds :

7-11 8-10 9-20 20-21

ring bonds :

1-2 1-6 1-9 2-3 3-4 4-5 5-6 6-7 7-8 8-9 11-12 11-15 12-13 13-14 14-15
14-16 15-19 16-17 17-18 18-19 21-22 21-26 22-23 23-24 24-25 25-26

exact/norm bonds :

1-2 1-6 1-9 2-3 3-4 4-5 5-6 6-7 7-8 7-11 8-9 8-10 9-20 11-12 11-15
12-13 13-14 14-15 14-16 15-19 16-17 17-18 18-19 20-21 21-22 21-26 22-23
23-24 24-25 25-26

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
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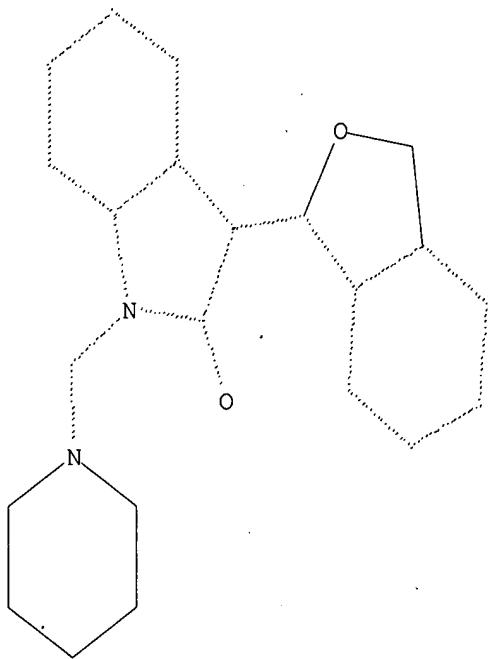
L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR

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Structure attributes must be viewed using STN Express query preparation.

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SAMPLE SCREEN SEARCH COMPLETED - 17 TO ITERATE

100.0% PROCESSED 17 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01
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BATCH **COMPLETE**
PROJECTED ITERATIONS: 93 TO 587
PROJECTED ANSWERS: 0 TO 0
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L2 0 SEA SSS SAM L1

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SEARCH TIME: 00.00.01
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L3 3 SEA SSS FUL L1

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                           ENTRY SESSION
FULL ESTIMATED COST           172.55 172.76
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FILE COVERS 1907 - 3 Jan 2007 VOL 146 ISS 2
FILE LAST UPDATED: 2 Jan 2007 (20070102/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13
L4 2 L3

=> d ed ibib abs hitstr 1-2

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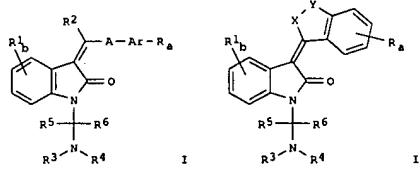
L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
 ED Entered STN: 28 May 2004
 ACESSION NUMBER: 2004:433773 HCAPLUS
 DOCUMENT NUMBER: 140:423586
 TITLE: Preparation of dihydroindolones as tyrosine kinase inhibitors for the treatment of disease
 INVENTOR(S): Andrews, Steven W.; Garst, Michael E.; Guo, Xialing; Hebert, Jonathan J.; Malone, Thomas; Wurster, Julie A.; Hull, Clarence Eugene
 PATENT ASSIGNEE(S): Allergan, Inc., USA
 SOURCE: U.S. Pat. Appl. Publ., 18 pp., Cont. of U.S. Ser. No. 306,975, abandoned.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PARENT -

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|-------------|
| US 2004102509 | A1 | 20040527 | US 2003-389416 | 20030313 |
| US 6747025 | B2 | 20040608 | | |
| CA 2507780 | A1 | 20040617 | CA 2003-2507780 | 20031119 |
| WO 2004050621 | A2 | 20040617 | WO 2003-US36988 | 20031119 |
| WO 2004050621 | A3 | 20040715 | | |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| AU 2003295658 | A1 | 20040623 | AU 2003-295658 | 20031119 |
| GB 2410744 | A | 20050810 | GB 2005-11267 | 20031119 |
| GB 2410744 | B | 20060412 | | |
| BR 2003016744 | A | 20051018 | BR 2003-16744 | 20031119 |
| JP 2006512400 | T | 20060413 | JP 2004-570761 | 20031119 |
| US 2004198802 | A1 | 20041007 | US 2004-829139 | 20040420 |
| PRIORITY APPLN. INFO.: | | | US 2003-306975 | B1 20021127 |
| | | | | |
| | | | US 2002-307097 | A 20021127 |
| | | | US 2003-389416 | A 20030313 |
| | | | WO 2003-US36988 | W 20031119 |

OTHER SOURCE(S): MARPAT 140:423586
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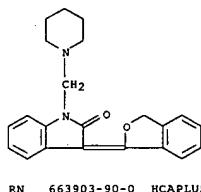
L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)



AB The title compds. (I or II; X = O, C(R2)2; Y = [C(R2)2]c; A = NR2, absent; R1 = halo, OH, NO2, CN, etc.; R2 = H, alkyl, Ph, etc.; R = halo, (un)substituted hydrocarbyl; R3, R4 = H, (un)substituted hydrocarbyl; NR3R4 = (heterocyclic ring); R5, R6 = H, alkyl, aryl; a = 0-3; b = 0-3; c = 1-2) which are capable of modulating tyrosine kinase signal transduction in order to regulate, modulate and/or inhibit abnormal cell proliferation, were prepared. Thus, reacting 4-morpholinaniline with 3-(hydroxymethylene)-3-dihydroindol-2-one (preparation given) in THF afforded 92% 3-[4-morpholinophenylamino]-methylene]-1,3-dihydroindol-2-one which showed IC50 of 260 nM against VEGFR2 kinase.

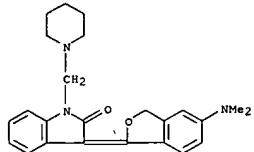
IT 663903-86-4 663903-90-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); B101 (Biological study); PREP (Preparation); USES (Uses)
 (preparation of dihydroindolones as tyrosine kinase inhibitors for treatment of disease)

RN 663903-86-4 HCAPLUS
 CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



RN 663903-90-0 HCAPLUS

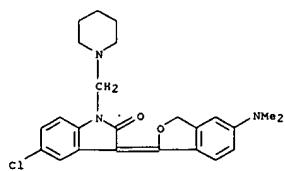
L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)
 CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of dihydroindolones as tyrosine kinase inhibitors for treatment of disease)

RN 663903-91-1 HCAPLUS

CN 2H-Indol-2-one, 5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
 ED Entered STN: 04 Mar 2004
 ACESSION NUMBER: 2004:176559 HCAPLUS
 DOCUMENT NUMBER: 140:210752
 TITLE: Dihydroindolone compound tyrosine kinase inhibitors for the treatment of disease
 INVENTOR(S): Andrews, Steven W.; Hebert, Jonathan J.
 PATENT ASSIGNEE(S): Allergan, Inc., USA
 SOURCE: U.S., 14 pp.
 CODEN: USXXAM

DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|------------|
| US 6699863 | B1 | 20040302 | US 2002-307097 | 20021127 |
| CA 2507780 | A1 | 20040617 | CA 2003-2507780 | 20031119 |
| WO 2004050621 | A2 | 20040617 | WO 2003-US36988 | 20031119 |
| WO 2004050621 | A3 | 20040715 | | |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| AU 2003295658 | A1 | 20040623 | AU 2003-295658 | 20031119 |
| GB 2410744 | A | 20050810 | GB 2005-11267 | 20031119 |
| GB 2410744 | B | 20060412 | | |
| BR 2003016744 | A | 20051018 | BR 2003-16744 | 20031119 |
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| PRIORITY APPLN. INFO.: | | | US 2002-306975 | A 20021127 |
| | | | US 2002-307097 | A 20021127 |
| | | | US 2003-389416 | A 20030313 |
| | | | WO 2003-US36988 | W 20031119 |

OTHER SOURCE(S): MARPAT 140:210752

AB The invention discloses organic mol.s., especially dihydroindolone derivs. (preparation

described) capable of modulating tyrosine kinase signal transduction in order to regulate, modulate and/or inhibit abnormal cell proliferation.

IT 663903-86-4P 663903-90-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); B101 (Biological study); PREP (Preparation); USES (Uses)

(dihydroindolone derivative tyrosine kinase inhibitors for treatment of disease)

RN 663903-86-4 HCAPLUS

CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)

AB M

unassigned

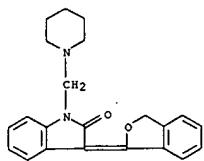
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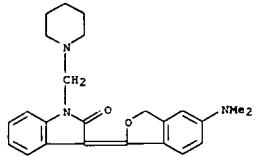
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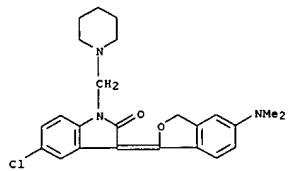
L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)



RN 663903-90-0 HCAPLUS
CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(dihydroindolone derivative tyrosine kinase inhibitors for treatment
of
disease)
RN 663903-91-1 HCAPLUS
CN 2H-Indol-2-one,
5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN (Continued)

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L1 STRUCTURE uploaded
L2 0 S L1
L3 3 S L1 FULL

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L4 2 S L3

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| COST IN U.S. DOLLARS | SINCE FILE ENTRY | TOTAL SESSION |
|--|------------------|---------------|
| FULL ESTIMATED COST | 15.74 | 188.50 |
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE ENTRY | TOTAL SESSION |
| CA SUBSCRIBER PRICE | -1.56 | -1.56 |

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NEWS 23 DEC 11 CAS REGISTRY chemical nomenclature enhanced
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NEWS X25 X.25 communication option no longer available

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DICTIONARY FILE UPDATES: 2 JAN 2007 HIGHEST RN 916646-22-5

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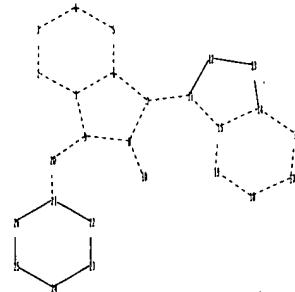
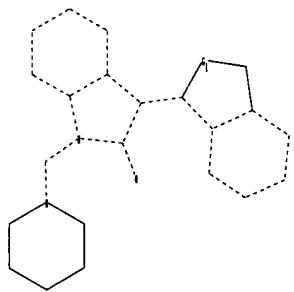
Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/reqprops.html>

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10829139s2



chain nodes :

10 20

ring nodes :

1 2 3 4 5 6 7 8 9 11 12 13 14 15 16 17 18 19 21 22 23 24 25
26

chain bonds :

7-11 8-10 9-20 20-21

ring bonds :

1-2 1-6 1-9 2-3 3-4 4-5 5-6 6-7 7-8 8-9 11-12 11-15 12-13 13-14 14-15
14-16 15-19 16-17 17-18 18-19 21-22 21-26 22-23 23-24 24-25 25-26

exact/norm bonds :

1-2 1-6 1-9 2-3 3-4 4-5 5-6 6-7 7-8 7-11 8-9 8-10 9-20 11-12 11-15
12-13 13-14 14-15 14-16 15-19 16-17 17-18 18-19 20-21 21-22 21-26 22-23
23-24 24-25 25-26

G1:C,O

Match level :

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11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
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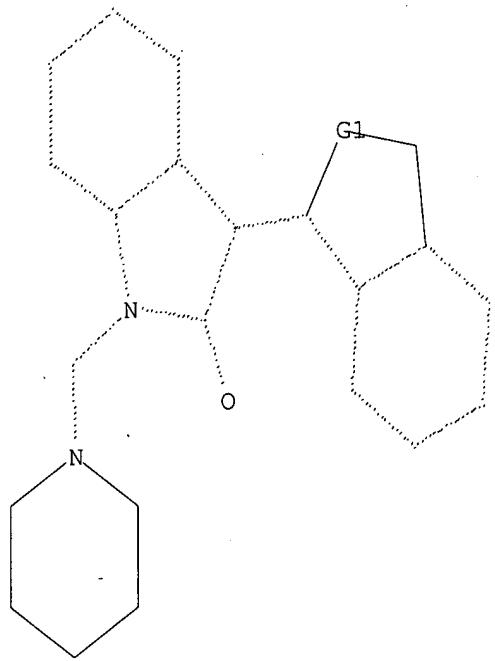
L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR

10829139s2



G1 C,O

Structure attributes must be viewed using STN Express query preparation.

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100.0% PROCESSED 36 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 360 TO 1080
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s 11 full
FULL SEARCH INITIATED 18:32:49 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 629 TO ITERATE

100.0% PROCESSED 629 ITERATIONS 3 ANSWERS
SEARCH TIME: 00.00.01

L3 3 SEA SSS FUL L1

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COST IN U.S. DOLLARS SINCE FILE TOTAL
FULL ESTIMATED COST ENTRY SESSION
172.10 172.31

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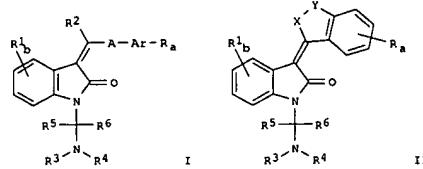
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L4 ANSWER 1 OF 2 HCPLUS COPYRIGHT 2007 ACS on STN
 ED Entered STN: 28 May 2004
 ACESSION NUMBER: 2004:433773 HCPLUS
 DOCUMENT NUMBER: 140:423586
 TITLE: Preparation of dihydroindolones as tyrosine kinase inhibitors for the treatment of disease
 INVENTOR(S): Andrews, Steven W.; Garst, Michael E.; Guo, Xialing; Hebert, Jonathan J.; Malone, Thomas; Wurster, Julie A.; Hull, Clarence Eugene
 PATENT ASSIGNEE(S): Allergan, Inc., USA
 SOURCE: U.S. Pat. Appl. Publ., 18 pp., Cont. of U.S. Ser. No. 306,975, abandoned.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|-------------|
| US 2004102509 | A1 | 20040527 | US 2003-389416 | 20030313 |
| US 6747025 | B2 | 20040608 | | |
| CA 2507780 | A1 | 20040617 | CA 2003-2507780 | 20031119 |
| WO 2004050621 | A2 | 20040617 | WO 2003-US36988 | 20031119 |
| WO 2004050621 | A3 | 20040715 | | |
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| AU 2003295658 | A1 | 20040623 | AU 2003-295658 | 20031119 |
| GB 2410744 | A | 20050810 | GB 2005-11267 | 20031119 |
| GB 2410744 | B | 20060412 | | |
| BR 2003016744 | A | 20051018 | BR 2003-16744 | 20031119 |
| JP 2006512400 | T | 20060413 | JP 2004-570761 | 20031119 |
| US 2004198802 | A1 | 20041007 | US 2004-829139 | 20040420 |
| PRIORITY APPLN. INFO.: | | | US 2002-306975 | B1 20021127 |
| | | | US 2002-307097 | A 20021127 |
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| | | | WO 2003-US36988 | W 20031119 |

OTHER SOURCE(S): MARPAT 140:423586
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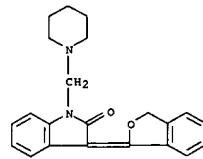
L4 ANSWER 1 OF 2 HCPLUS COPYRIGHT 2007 ACS on STN (Continued)



AB The title compds. [I or II; X = O, C(R2)2; Y = [C(R2)2]c; A = NR2, absent; R1 = halo, OH, NO2, CN, etc.; R2 = H, alkyl, Ph, etc.; R = halo, (un)substituted hydrocarbyl; R3, R4 = (hetero)cyclic ring; R5, R6 = H, alkyl, aryl; a = 0-3; b = 0-3; c = 1-2] which are capable of modulating tyrosine kinase signal transduction in order to regulate, modulate and/or inhibit abnormal cell proliferation, were prepared. Thus, reacting 4-morpholinoaniline with 3-(hydroxymethylene)-3-(4-dihydroindol-2-one (preparation given) in THF afforded 92% 3-[4-(morpholino)phenylamino]-1,3-dihydroindol-2-one which showed IC50 of 260 nM against VEGFR2 kinase.

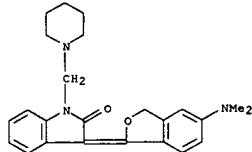
IT 663903-86-4P 663903-90-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of dihydroindolones as tyrosine kinase inhibitors for treatment of disease)

RN 663903-86-4 HCPLUS
 CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)

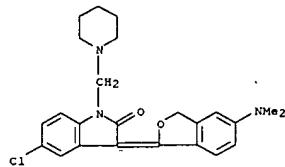


RN 663903-90-0 HCPLUS

L4 ANSWER 1 OF 2 HCPLUS COPYRIGHT 2007 ACS on STN (Continued)
 CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of dihydroindolones as tyrosine kinase inhibitors for treatment of disease)
 RN 663903-91-1 HCPLUS
 CN 2H-Indol-2-one, 5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 HCPLUS COPYRIGHT 2007 ACS on STN
 ED Entered STN: 04 Mar 2004
 ACESSION NUMBER: 2004:176559 HCPLUS
 DOCUMENT NUMBER: 140:210752
 TITLE: Dihydroindolone compound tyrosine kinase inhibitors for the treatment of disease
 INVENTOR(S): Andrews, Steven W.; Hebert, Jonathan J.
 PATENT ASSIGNEE(S): Allergan, Inc., USA
 SOURCE: U.S., 14 pp.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

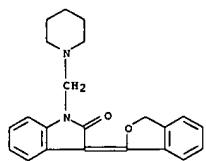
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| US 6699863 | B1 | 20040302 | US 2002-307097 | 20021127 |
| CA 2507780 | A1 | 20040617 | CA 2003-2507780 | 20031119 |
| WO 2004050621 | A2 | 20040617 | WO 2003-US36988 | 20031119 |
| WO 2004050621 | A3 | 20040715 | | |
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| AU 2003295658 | A1 | 20040623 | AU 2003-295658 | 20031119 |
| GB 2410744 | A | 20050810 | GB 2005-11267 | 20031119 |
| GB 2410744 | B | 20060412 | | |
| BR 2003016744 | A | 20051018 | BR 2003-16744 | 20031119 |
| JP 2006512400 | T | 20060413 | JP 2004-570761 | 20031119 |
| PRIORITY APPLN. INFO.: | | | US 2002-306975 | A 20021127 |
| | | | US 2002-307097 | A 20021127 |
| | | | US 2003-389416 | A 20030313 |
| | | | WO 2003-US36988 | W 20031119 |

OTHER SOURCE(S): MARPAT 140:210752
 AB The invention discloses organic mol., especially dihydroindolone derivs., (preparation described) capable of modulating tyrosine kinase signal transduction in order to regulate, modulate and/or inhibit abnormal cell proliferation.
 IT 663903-86-4P 663903-90-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (dihydroindolone derivative tyrosine kinase inhibitors for treatment of disease)

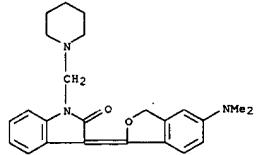
RN 663903-86-4 HCPLUS
 CN 2H-Indol-2-one, 1,3-dihydro-3-(1(3H)-isobenzofuranylidene)-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)

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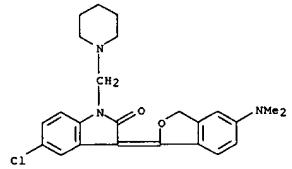
L4 ANSWER 2 OF 2 HCPLUS COPYRIGHT 2007 ACS on STN (Continued)



RN 663903-90-0 HCPLUS
CN 2H-Indol-2-one, 3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



IT 663903-91-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(dihydroindolone derivative tyrosine kinase inhibitors for treatment
of
disease)
RN 663903-91-1 HCPLUS
CN 2H-Indol-2-one,
5-chloro-3-[5-(dimethylamino)-1(3H)-isobenzofuranylidene]-1,3-dihydro-1-(1-piperidinylmethyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 2 HCPLUS COPYRIGHT 2007 ACS on STN (Continued)

REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR
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| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE | TOTAL |
| | ENTRY | SESSION |
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